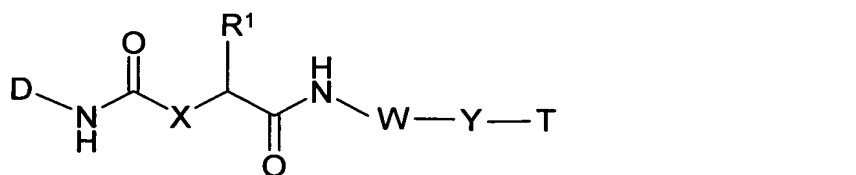


This listing of claims will replace all prior versions of claims in the application.

Listing of Claims: Please amend the claims as follows:

We claim:

Claim 1. (Currently Amended) ~~Compounds~~ A compound of the formula I



in which wherein

D denotes is an aromatic five-membered heterocyclic ring having 1 to 4 N, O and/or S atoms which is unsubstituted or mono- or polysubstituted by Hal, A, OR², N(R²)₂, NO₂, CN, COOR² or CON(R²)₂,

X denotes is NR³ or O,

R¹ denotes is H, Ar, Het, cycloalkyl or A, which ~~may be~~ is optionally substituted by OR², SR², N(R²)₂, Ar, Het, cycloalkyl, CN, COOR² or CON(R²)₂,

R² denotes is H, A, -[C(R³)₂]_n-Ar, -[C(R³)₂]_n-Het, -[C(R³)₂]_n-cycloalkyl, -[C(R³)₂]_n-N(R³)₂ or -[C(R³)₂]_n-OR³,

R³ denotes is H or A,

W denotes is -[C(R³)₂]_n-,

Y denotes is alkylene, cycloalkylene, Het-diyl or Ar-diyl,

T denotes is a mono- or bicyclic saturated, unsaturated or aromatic carbo- or heterocyclic ring having 0 to 4 N, O and/or S atoms, which ~~may be~~ is optionally unsubstituted or mono-, di- or trisubstituted by Hal, A, -[C(R³)₂]_n-Ar, -[C(R³)₂]_n-Het, -[C(R³)₂]_n-cycloalkyl, OR³, N(R³)₂, NO₂, CN, COOR², CON(R²)₂, NR²COA, NR²CON(R²)₂, NR²SO₂A, COR², SO₂NR² and/or S(O)_mA and/or carbonyl oxygen,

or $N(R^2)_2$

and, if Y = piperidine-1,4-diyl, also R^2 or cycloalkyl,

A denotes is unbranched or branched alkyl having 1-10 C atoms, in which wherein one or two CH_2 groups ~~may be~~ are optionally replaced by O or S atoms and/or by $-CH=CH-$ groups and/or ~~also~~ wherein 1-7 H atoms ~~may be~~ are optionally replaced by F,

Ar denotes is phenyl, naphthyl or biphenyl, each of which is, independently of one another, unsubstituted or mono-, di- or trisubstituted by Hal, A, OR^3 , $N(R^3)_2$, NO_2 , CN, $COOR^3$, $CON(R^3)_2$, NR^3COA , $NR^3CON(R^3)_2$, NR^3SO_2A , COR^3 , $SO_2N(R^3)_2$, $S(O)_mA$, $-[C(R^3)_2]_n-COOR^{2'}$ or $-O-[C(R^3)_2]_o-COOR^{2'}$,

$R^{2'}$ denotes is H, A, $-[C(R^3)_2]_n-Ar'$, $-[C(R^3)_2]_n-Het'$, $-[C(R^3)_2]_n-cycloalkyl$, $-[C(R^3)_2]_n-N(R^3)_2$ or $-[C(R^3)_2]_n-OR^3$,

$R^{2''}$ denotes is H, A, $-[C(R^3)_2]_n-Ar'$ or $-[C(R^3)_2]_n-cycloalkyl$, $-[C(R^3)_2]_n-N(R^3)_2$ or $-[C(R^3)_2]_n-OR^3$,

Ar' denotes is phenyl or benzyl, each of which is, independently of one another, unsubstituted or mono- or disubstituted by Hal or A,

Het denotes is a mono- or bicyclic saturated, unsaturated or aromatic heterocyclic ring having 1 to 4 N, O and/or S atoms, which ~~may be~~ is unsubstituted or mono-, di- or trisubstituted by carbonyl oxygen, $=S$, $=N(R^3)_2$, Hal, A, $-[C(R^3)_2]_n-Ar$, $-[C(R^3)_2]_n-Het^1$, $-[C(R^3)_2]_n-cycloalkyl$, $-[C(R^3)_2]_n-OR^{2'}$, $-[C(R^3)_2]_n-N(R^2)_2$, NO_2 , CN, $-[C(R^3)_2]_n-COOR^{2'}$, $-[C(R^3)_2]_n-CON(R^2)_2$, $-[C(R^3)_2]_n-NR^{2'}COA$, $NR^{2'}CON(R^2)_2$, $-[C(R^3)_2]_n-NR^{2'}SO_2A$, $COR^{2'}$, $SO_2NR^{2'}$ and/or $S(O)_mA$,

Het^1 denotes is a mono- or bicyclic saturated, unsaturated or aromatic heterocyclic ring having 1 to 2 N, O and/or S atoms, which ~~may be~~ is unsubstituted or mono- or disubstituted by carbonyl oxygen, $=S$, $=N(R^3)_2$, Hal, A, $OR^{2''}$, $N(R^{2''})_2$, NO_2 , CN, $COOR^{2''}$, $CON(R^{2''})_2$, $NR^{2''}COA$, $NR^{2''}CON(R^{2''})_2$, $NR^{2''}SO_2A$, $COR^{2''}$, $SO_2NR^{2''}$ and/or $S(O)_mA$,

Hal denotes is F, Cl, Br or I,

n denotes is 0, 1 or 2,

m denotes is 0, 1 or 2,

o denotes is 1, 2 or 3,
and or a pharmaceutically usable derivatives acceptable salt, solvate[s]
and or stereoisomer[s] thereof, including or a mixture[s] thereof in all
ratios.

Claim 2. (Currently Amended) Compounds A compound according to Claim 1, in
which wherein

D denotes is an aromatic five-membered heterocyclic ring having 1
to 2 N, O and/or S atoms which is unsubstituted or mono- or
disubstituted by Hal,
and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or
stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 3. (Currently Amended) Compounds A compound according to Claim 1, in
which wherein

D denotes is a thienyl ring which is mono- or disubstituted by Hal,
and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or
stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 4. (Currently Amended) Compounds A compound according to Claim 1, in
which wherein

R² denotes is H or alkyl having 1, 2, 3, 4, 5 or 6 C atoms,
and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or
stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 5. (Currently Amended) Compounds A compound according to Claim 1, in
which wherein

R¹ denotes is H or unsubstituted phenyl, thienyl or alkyl having 1-6
C atoms,
and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or
stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 6. (Currently Amended) Compounds A compound according to Claim 1, in which wherein

X denotes is NH or O,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 7. (Currently Amended) Compounds A compound according to Claim 1, in which wherein

W denotes is (CH₂)_n,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 8. (Currently Amended) Compounds A compound according to Claim 1, in which wherein

Y denotes is Ar-diyl or Het-diyl,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 9. (Currently Amended) Compounds A compound according to Claim 1, in which wherein

T denotes is a mono- or bicyclic saturated, unsaturated or aromatic heterocyclic ring having 1 to 2 N and/or O atoms, which ~~may be~~ is unsubstituted or mono- or disubstituted by carbonyl oxygen, or N(R²)₂

and, if Y = piperidine-1,4-diyl, also R²,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 10. (Currently Amended) Compounds A compound according to Claim 1, in which wherein

T denotes is a mono- or bicyclic saturated or unsaturated heterocyclic ring having 1 to 2 N and/or O atoms which is mono- or

disubstituted by carbonyl oxygen (=O),

or $N(R^2)_2$

and, if Y = piperidine-1,4-diyl, also R^2 ,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 11. (Currently Amended)

Compounds A compound according to Claim 1,

in which wherein

T denotes is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl, each of which is mono- or disubstituted by carbonyl oxygen, or $N(R^2)_2$

and, if Y = piperidine-1,4-diyl, also R^2 ,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 12. (Currently Amended)

Compounds A compound according to Claim 1,

in which wherein

Ar denotes is phenyl which is unsubstituted or mono- or disubstituted by Hal, A, OA, SO_2A , $COOR^2$, SO_2NH_2 or CN,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 13. (Currently Amended)

Compounds A compound according to Claim 1,

in which wherein

D denotes is an aromatic five-membered heterocyclic ring having 1 to 2 N, O and/or S atoms which is unsubstituted or mono- or disubstituted by Hal,

R^1 denotes is H or unsubstituted phenyl, thienyl or alkyl having 1-6 C atoms,

R^2 denotes is H or alkyl having 1, 2, 3, 4, 5 or 6 C atoms,

X denotes NH or O,
W denotes is W (CH₂)_n,
Y denotes is Ar-diyl, pyridinediyl or piperidinediyl,
Ar denotes is phenyl which is unsubstituted or mono- or disubstituted by Hal, A, OA, SO₂A, COOR², SO₂NH₂ or CN,
T denotes is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl, each of which is mono- or disubstituted by carbonyl oxygen, or N(R²)₂
and, if Y = piperidine-1,4-diyl, also R²,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, ~~including or a mixture[s]~~ thereof in all ratios.

Claim 14. (Currently Amended) ~~Compounds~~ A compound according to Claim 1, ~~in which~~ wherein

D denotes is thienyl, thiazolyl or furyl, each of which is mono- or disubstituted by Hal,
R¹ denotes is H or unsubstituted phenyl, thienyl or alkyl having 1-6 C atoms,
R² denotes is H or alkyl having 1, 2, 3, 4, 5 or 6 C atoms,
X denotes is NH or O,
W denotes is W (CH₂)_n,
Y denotes is Ar-diyl, pyridinediyl or piperidinediyl,
Ar denotes is phenyl which is unsubstituted or mono- or disubstituted by Hal, A, OA, SO₂A, COOR², SO₂NH₂ or CN,
T denotes is piperidin-1-yl, pyrrolidin-1-yl, pyridinyl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, pyridazin-2-yl, pyrazinyl, azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl, each of which is unsubstituted or mono- or disubstituted by carbonyl oxygen, or N(R²)₂
and, if Y = piperidine-1,4-diyl, also R²,

and or a pharmaceutically usable ~~derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, ~~including or a~~ mixture[s] thereof in all ratios.

Claim 15. (Currently Amended) ~~Compounds~~ A compound according to Claim 1 selected from the group which is

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]valeramide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-3-methylphenyl]valeramide,

2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]acetamide,

(R)-2-[3-(5-bromothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]valeramide,

(R)-2-[3-(5-bromofuran-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]valeramide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]-2-phenylacetamide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]-2-(thiophen-2-yl)acetamide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(2-oxo-1*H*-pyrazin-1-yl)-phenyl]valeramide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[2-oxo-3,4,5,6-tetrahydro-[1,2']bipyridinyl-5'-yl]valeramide,

(S)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]-2-phenylacetamide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenylmethyl]valeramide,

(R)-2-[3-(5-chlorothiazol-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]valeramide,

(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-[[4-(3-oxo-morpholin-4-yl)phenyl]valeramide,

(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-[C-(3,4,5,6-tetrahydro-2H-[1,4']bipyridinyl-4-yl)methyl]valeramide,

(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-[1-isopropyl-piperidin-4-ylmethyl]-2-phenylacetamide,

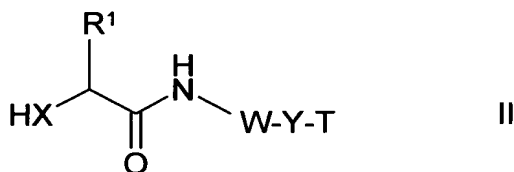
(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-[[4-(morpholin-4-yl)phenyl]valeramide; or

(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-(4-dimethylamino-phenyl)-2-phenylacetamide,

and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

Claim 16. (Currently Amended) ~~Process A~~ process for the preparation of compounds a compound of the formula I according to Claim 1 and pharmaceutically usable derivatives, solvates and stereoisomers thereof, characterised in that comprising

a) reacting a compound of the formula II



in which wherein

R¹, W, X, Y and T have the meaning indicated in Claim 1,

~~is reacted~~ with a compound of the formula III



in which wherein

D has the meaning indicated in Claim 1,

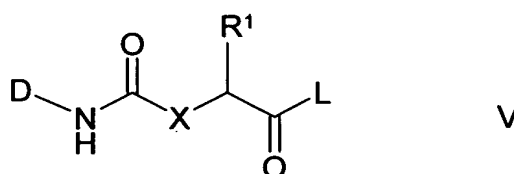
or

b) reacting a compound of the formula IV



~~in which~~ wherein W, Y and T have the meaning indicated in Claim 1,

~~is reacted~~ with a compound of the formula V



~~in which~~ wherein

L denotes Cl, Br, I or a free or reactively functionally modified OH group, and

R¹, X and D have the meanings indicated in Claim 1,
and/or

optionally converting a base or acid of the formula I ~~is converted~~ into one of its salts.

Claim 17. (Currently Amended) ~~Compounds of the formula I according to Claim 1 as inhibitors of~~ A method of inhibiting coagulation factor Xa comprising contacting said coagulation factor Xa with a compound according to claim 1.

Claim 18. (Currently Amended) ~~Compounds of the formula I according to Claim 1 as inhibitors of~~ A method of inhibiting coagulation factor VIIa comprising contacting said coagulation factor VIIa with a compound according to claim 1.

Claim 19. (Currently Amended) ~~Medicaments~~ A pharmaceutical composition comprising at least one compound of the formula I according to Claim 1 ~~and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and optionally excipients and/or adjuvants~~ and a pharmaceutically

acceptable carrier.

Claim 20. (Currently Amended) ~~Medicaments~~ A pharmaceutical composition comprising at least one compound ~~of the formula-I according to Claim 1 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and at least one further medicament active ingredient~~ an excipient, adjuvant, or vitamin.

Claim 21. (Withdrawn-Currently Amended) ~~Use of compounds according to Claim 1 and/or physiologically acceptable salts and solvates thereof for the preparation of a medicament~~ A method for the treatment of thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, tumours, tumour diseases and/or tumour metastases comprising administering to a subject in need thereof a compound of claim 1.

Claim 22. (Currently Amended) ~~Set (kit) consisting of separate packs of~~ A set or a kit comprising

- (a) an effective amount of a compound ~~of the formula-I according to Claim 1 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios,~~
- and
- (b) an effective amount of ~~a further medicament active ingredient~~ an excipient, adjuvant, or vitamin.

Claim 23. (Withdrawn) ~~Use of compounds according to Claim 1 and/or physiologically acceptable salts and solvates thereof for the preparation of a medicament~~ A method for the treatment of thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, tumours, tumour diseases and/or tumour metastases comprising administering to a subject in need thereof a pharmaceutical composition of claim 19. , in combination with at least one further medicament active ingredient.